## PATENT COOPERATION TREATY

# **PCT**

#### INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference 15270C-4-2PC	FOR FURTHER ACTION	see Form PCT/ISA/220 as well as, where applicable, item 5 below.
International application No. PCT/US 08/80382	International filing date (day/mod 17 October 2008 (17.10.2008)	nth/year) (Earliest) Priority Date (day/month/year) 17 October 2007 (17.10.2007)
Applicant ELAN PHARMA INTERNATIONAL LIMITE	ED .	
according to Article 18. A copy is bein  This international search report consists	ig transmitted to the International B	
a translation of the international search authorized by or notified to c. With regard to any nucleon.  Certain claims were four to the international search authorized by or notified to c. With regard to any nucleon.  Unity of invention is lack to the title, the text is approved as subtractions.	blication in the language in which in international application into ed for the purposes of international report has been established taking to this Authority under Rule 91 (Ru bide and/or amino acid sequence and unsearchable (see Box No. II).	which is the language of search (Rules 12.3(a) and 23.1(b)).  into account the rectification of an obvious mistake le 43.6bis(a)).  disclosed in the international application, see Box No. I.
may, within one month from  6. With regard to the <b>drawings</b> ,  a. the figure of the <b>drawings</b> to be as suggested by the as selected by this A as selected by this A	ed, according to Rule 38.2(b), by the second of the second	ed to suggest a figure.

Form PCT/ISA/210 (first sheet) (April 2007)

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Box	No. l	I	Nucleotide and/or amino acid sequence(s) (Continuation of item1.b of the first sheet)
1.	With carrie	regar ed out	d to any nucleotide and/or amino acid sequence disclosed in the international application, the international search was on the basis of:
i	a.	type o	f material a sequence listing table(s) related to the sequence listing
	b.	forma	t of material on paper in electronic form
	c.	time o	of filing/furnishing  contained in the international application as filed  filed together with the international application in electronic form  furnished subsequently to this Authority for the purposes of search
2.		or f	addition, in the case that more than one version or copy of a sequence listing and/or table relating thereto has been filed furnished, the required statements that the information in the subsequent or additional copies is identical to that in the lication as filed or does not go beyond the application as filed, as appropriate, were furnished.
3.	Addi	itional	comments:
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Box No. II	Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)			
This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:				
1. Cl be	aims Nos.: cause they relate to subject matter not required to be searched by this Authority, namely:			
be	aims Nos.: cause they relate to parts of the international application that do not comply with the prescribed requirements to such an tent that no meaningful international search can be carried out, specifically:			
3. Cl	aims Nos.: 4-9 and 61-63 cause they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).			
Box No. III	Observations where unity of invention is lacking (Continuation of item 3 of first sheet)			
	tional Searching Authority found multiple inventions in this international application, as follows:			
This applicat concept und	ion contains the following inventions or groups of inventions which are not so linked as to form a single general inventive er PCT Rule 13.1. In order for all inventions to be examined, the appropriate additional examination fees must be paid.			
	n 1-3, 10-60, and 64-133 are directed to a method of treating Alzheimer's disease, and related diseases relative to the ApoE4 alleles.			
Group II clai	ms 134-136 are directed to humanized form of a 10D5 antibody.			
Group III cla	ims 137-139 are directed to a humanized form of a 12A11 antibody.			
Group IV cla	ims 140-142 are directed to a humanized form of a 3D6 antibody.			
	**************************************			
	s all required additional search fees were timely paid by the applicant, this international search report covers all searchable aims.			
	s all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of ditional fees.			
on on	s only some of the required additional search fees were timely paid by the applicant, this international search report covers ly those claims for which fees were paid, specifically claims Nos.: 3, 10-60, 64-133, 143, and 145-146			
4. No res	o required additional search fees were timely paid by the applicant. Consequently, this international search report is stricted to the invention first mentioned in the claims; it is covered by claims Nos.:			
Remark on	The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.  The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.  No protest accompanied the payment of additional search fees.			

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A. CLASSIFICATION OF SUBJECT MATTER  IPC(8) - A61K 39/00; C07K 16/18 (2009.01)  USPC - 424/133.1, 530/387.3  According to International Patent Classification (IPC) or to both national classification and IPC					
	DS SEARCHED		1-7-7-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1		
Minimum documentation searched (classification system followed by classification symbols) IPC(8) - A61K 39/00; C07K 16/18 (2009.01) USPC - 424/133.1, 530/387.3; 424/130.1, 424/141.1, 530/387.3, 530/350, 530/300					
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched IPC(8) - A61K 39/00; C07K 16/18 (2009.01) - see keyword below USPC - 424/133.1, 530/387.3; 424/130.1, 424/141.1, 530/387.3, 530/350, 530/300 - see keyword below					
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) PubWEST(USPT,PGPB,EPAB,JPAB); Medline, Google Search terms: Alzheimer's ApoE4 alleles, ApoE4 non-carrier, apolipoprotein E, zero, beta-amyloid, antibody, N-terminal, epitope, mg/kg, intravenous, infusion, ?g/ml, 3D6 antibody, PTA-5130, bapineuzumab, pg/ml, plasma concentration, PTA-5130, vasogenic edema					
C. DOCUM	TENTS CONSIDERED TO BE RELEVANT				
Category*	Citation of document, with indication, where ap	ppropriate, of the relevant passages	Relevant to claim No.		
l i	US 2006/0280743 A1 (BASI et al.) 14 December 2006 [0057], [0074], [0099], [0105], [0114], [0171], [0172], [0	174], [0199], [0201], [0204], [0208],	1-3, 10-13, 15-21, 23-25, 27-28, 30-32, 97-98, 101-103, 111-116, 121, 128-131, 133, 145-146		
	io209j, io215j, io216j, io217j, io2210j, io232j, io238j, l io363j	(U248], [U274], [U339], [U337], [U39U], ANU	14, 22, 26, 29, 33-60, 64- 87, 89-95, 99, 104, 106- 108, 110, 117-119, 123- 127, 132		
			88, 96, 100, 105, 109, 120, 122, 143		
Y	US 2007/0196375 A1 (TOBINICK) 23 August 2007 (2	3.08.2007), para [0019], and [0267]	14, 22, 26, 48, 60, 66-70, 87, 94-95, 99, 104, 108		
	Kinnecom et. al. Course of cerebral amyloid angiopath April. Vol. 68(17), p. 1411-6. Abstract; pg 1411, para 2 included to establish publication date)	y?related inflammation. Neurology. 2007 and 3; pg 1415, col 1, last para (abstract	29, 33-60, 64-87, 89-95, 106-108, 110, 117-119, 123-127, 132		
			88, 96, 100, 105, 109, 120, 122, 143		
Further documents are listed in the continuation of Box C.					
* Special categories of cited documents:  "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand					
to be of particular relevance the principle or theory underlying the invention  "E" earlier application or patent but published on or after the international filing date "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive					
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)  step when the document is taken alone document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is					
"O" document referring to an oral disclosure, use, exhibition or other means combined with one or more other such documents, such combination being obvious to a person skilled in the art					
the priori	t published prior to the international filing date but later than ity date claimed		-		
Date of the actual completion of the international search  Date of mailing of the international search report  Standard 2010					
08 March 2009 (08.03.2009) 2 5 MAR 2009			<b></b>		
Name and mailing address of the ISA/US  Authorized officer:  Lee W. Young					
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Raccimile No.	571-273-3201	DOT OSD: 571-272-7774			

International application No.
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C (Continua	tion). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevan	Relevant to claim No.	
A	US 2006/0193850 A1 (WARNE et al.) 31 August 2006 (31.08.2006), para [0149]; SEQ ID NO: 1 and 2		88, 96, 100, 105, 109, 120, 122, 143
A	Aylward et al. Cerebellar volume in adults with Down syndrome. Arch Neurol. 1997 Feb;54(2):209-12. Abstract		1-3, 10-60, 64-133, 143 145-146
A	Cofke et al. Remifentanil-Induced Cerebral Blood Flow Effects in Normal Humans: Dose and ApoE Genotype. Neurosurg Anesthes Neurosci. July 2007, Vol. 105(1),p.167-175.		1-3, 10-60, 64-133, 143 145-146
	· 		

Form PCT/ISA/210 (continuation of second sheet) (April 2007)

International application No. PCT/US 08/80382

Continuation of

Box No. III (unity of invention is lacking)

Group V claim 143 is directed to a humanized antibody comprising a humanized light chain having an amino acid sequence comprising SEQ 10 NO:48 and a humanized heavy chain having an amino acid sequence comprising SEQ ID NO:66 or SEQ ID NO:67.

Group VI claim 144 is directed to an isolated nucleic acid having a sequence comprising SEQ ID NO:68 provided that residues 1-57 encoding a signal sequence mayor may not be present.

Group VII claims 145-146 are directed to an isolated humanized antibody comprising a mature light chain variable region sequence of SEQ ID NO:2 and a mature heavy chain variable region sequence of SEQ ID NO:3, and a human heavy chain constant region of IgG isotype with L234A, L235A, and G237A mutations, wherein positions are numbered by the EU numbering system.

Group VIII claims 147-148 are directed to an isolated humanized form of a 12B4 antibody, wherein the 12B4 antibody is characterized by a light chain variable region sequence of SEQ ID NO:31, and heavy chain variable region sequence of SEQ ID NO:32, and a human heavy chain constant region of IgG isotype with L234A, L235A, and G237A mutations, wherein positions are numbered by the EU numbering system.

Group IX claims 149-151 are directed to a humanized form of a 266 antibody (ATCC accession number PTA6123) comprising a human heavy chain constant region with L234A, L235A and G237A mutations, wherein positions are numbered by the EU numbering system.

Group X claims 152-160 are directed to an isolated antibody comprising a human heavy chain constant region of isotype IgGI, wherein amino acids at positions 234, 235, and 237 (EU numbering) are each alanine.

Group XI claims 161- 193 are directed to a method and a kit for determining a regime for bapineuzumab administration.

Group XII claims 194-195 are directed to a method for improving the safety of bapineuzumab.

The shared technical feature of Groups I, XI and XII is identifying patients having no ApoE4 alleles exhibiting various brain diseases, including, inter alia, Alzheimer's disease. However, this is not an improvement over the prior art of US 5773220 A to DeKoskey et al. (30 June 1998) that specifically teaches identifying patients having no ApoE4 alleles exhibiting various brain diseases, including, inter alia, Alzheimer's disease (abstract, col 2 in 5-20). Groups II-X are directed to various polypeptide sequences and/or nucleic acid sequences that share no common technical feature with each other or with Groups I, XI and XII, and do not relate to a single general inventive concept because, under PCT Rule 13.2, the different nucleotides or polypeptides represented by the nucleic acid sequences or amino acid sequences are not common to one another but are different because they are composed of unique structural sequences.

inventive concept because, under PCT Rule 13.2, the different nucleotides or polypeptides represented by the nucleic acid sequences or amino acid sequences are not common to one another but are different because they are composed of unique structural sequences.

Note that Claim Nos. 4-9 and 61-63 have been found to be unsearchable under Article 17(2)(b) because of defects under Article 17(2)(a) and therefore have not been included with any invention.